Bot.

substituted aryl or heteroaryl with at least one substituent selected from the group consisting of lower alkyl, lower alkoxy, lower alkylthio, lower alkoxycarbonyl, lower alkylsulfonyl, halogen, CF_3 , $-OCF_3$, -OH, $-NO_2$, -CN, aryl, aryloxy, cycloalkly and heterocycloalkyl, X is $-(CH_2)_n$ -Z, Z is selected from the group consisting of a covalent bond, -NH-, =0 and =S, n is 0, 1 or 2, Y is oxygen or sulfur, R_1 is selected from the group consisting of hydrogen, -OH, halogen, lower alkyl and lower alkoxy, the alkyl and alkoxy being unsubstituted or substituted with at least one member of the group consisting of $-CF_3$, lower alkoxy, $-NH_2$ and mono- and di-lower alkylamino. R_{2a} and R_{2b} are individually selected from the group consisting of hydrogen, substituted or unsubstituted lower alkyl, substituted or unsubstituted or unsubsti

 R_{23} , $R_{21}-Z_{21}$ and $Z_{22}-R_{24}$,

 R_{22} and R_{23} are individually selected from the group consisting of hydrogen, lower alkyl, cycloalkyl, cycloalkylalkyl, aryl, aralkyl, heteroaryl, or heteroarylalkyl, alkylsulfonyl, cycloalkylsulfonyl, arylsulfonyl, lower alkoxycarbonyl, aryloxycarbonyl alkylcarbonyl, arylcarbonyl and cycloalkylcarbonyl, Z_{21} and Z_{22} are individually selected from the group consisting of oxygen, sulfur, -CO- and -O-CO-, R_{24} is selected from the group consisting of hydrogen, lower alkyl, cycloalkyl, cycloalkylalkyl, aryl, aralkyl, heteroaryl,

heteroarylalkyl, alkylsulfonyl, cycloalkylsulfonyl and arylsulfonyl, R_{21} is selected from the group consisting of hydrogen, lower alkyl, aryl and aralkyl, R_3 is selected from the group consisting of hydrogen, halogen, $-NO_2$, -CN, unsubstituted or substituted alkyl of 1 to 10 carbon atoms, unsubstituted or substituted lower alkyl, unsubstituted or substituted lower alkynyl, unsubstituted or substituted or substituted or substituted or substituted or substituted aryl, unsubstituted or substituted aryl, unsubstituted or substituted aralkyl, unsubstituted or substituted lower aryloxalkyl, unsubstituted or substituted heteroaryl, unsubstituted or substituted heteroarylalkyl and $-Z_{31}R_{31}$, the substituents being selected from the group consisting of halogen, aryl, R_{32}

-N and - Z_{32} - R_{34} , - Z_{31} is selected from the group R_{33}

consisting of -O-, -C(O)-, -OC(O)- and -S- R_{31} is selected from the group consisting of hydrogen, lower alkyl, aryl and lower aralkyl, R_{32} and R_{33} are individually selected from the group consisting of hydrogen, lower alkyl, aralkyl and alkylcarbonyl or together with the nitrogen form a heterocyloalkyl, Z_{32} is selected from the group consisting of oxygen, sulfur, -C(O)-, -S(O), -OCO- and -SO₂, R_{34} is selected from the group consisting of hydrogen, lower alkyl, aryl and lower aralkyl and its non-toxic, pharmaceutically acceptable salts sufficient to treat said pathological disorders.

Claim 3/, line 1, cancel the amendment and insert -- The method of claim 10 wherein--

The method of claim 10 wherein the compound is selected from the

B² group consisting of/-

Claim 9, line 2, change "1" to --10--

11. A compound of the formula

B³

$$R'_1$$
 R'_2a'
 R'_2b'

wherein W' is hydrogen or -C(Y')-X-R', R' is selected from the group consisting of phenyl, naphthyl indolyl and pyridyl, all unsubstituted or substituted with at least one member of the group consisting of methyl, ethyl, propyl, isopropyl, butyl, tert.-butyl, methoxy, ethoxy, methylthio, ethylthio, methoxycarbonyl, ethoxycarbonyl, methylsulfonyl, ethylsulfonyl, chlorine, fluorine, bromine, trifluoromethyl, trifluoromethoxy, -OH, $-NO_2$, -CN, phenyl, phenoxy and morpholino, X' is selected from the group consisting of $-CH_2-$, $-CH_2-CH_2-$, $-CH_2NH-$, -NH-, -O-, -S- and a covalent bond, Y' is oxygen or sulfur, R'₁ is at least one member of the group consisting of hydrogen, chlorine, methyl and methoxy, R'_{2a} and R'_{1b} are individually hydrogen or methyl, R'₃ is selected from the group

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consisting of hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, methoxyethyl, ethoxyethyl, dimethylaminoethyl, cyclohexyl methyl, phenyl, diphenyl, benzyl unsubstituted or substituted with -OH or methoxy, phenethyl, naphthylmethyl and indolylmethyl excluding the compounds of Formula II wherein a) W' is hydrogen, R'₁ is o-chlorine, R'_{2a} is hydrogen, R'_{2b} is hydrogen or methyl and R'_3 is methyl and b) wherein W' is -(Y)C-X'-R' and i) X'is -NH-, Y' is oxygen, R \searrow is o-chlorine, R' $_{2a}$ and R' $_{2b}$ are hydrogen, R', is methyl and R' is selected from the group consisting of 4tert.butyl-phenyl, 4-trifluoxomethyl-phenyl, 4-methoxy-phenyl, 3,4,5-trimethoxy-phenyl, 2,3-dichloro-phenyl, 2,3-difluoro-phenyl, 4-phenoxy-phenyl, pyridinyl and cyanophenyl or ii) X' is -NH-, Y' is sulfur, R'_1 is o-chloro, R'_{2a} and R'_{2b} are hydrogen, R'_3 is methyl and R' is selected from the group consisting of 4-tert.butylphenyl, 2,4-ditert.butyl-phenyl, 2-trifl\u00e4oromethyl-phenyl, 3trifluoromethyl-phenyl, 4-trifluoromethyl-phenyl, 4-methoxy-phenyl, 3,4,5-trimethoxy-phenyl, 4-fluoro-phenyl and 4-methylsulfonylphenyl or iii) X' is -CH2-NH-, Y is oxygen, R'1 is &-chlorine, R'2a and R'_{2b} are hydrogen, R'_{3} is methyl and R' is phenyl, or iiii) X'is oxygen or a covalent bond, Y' is oxygen, R', is o-chlòrine, R'_{2a} and R'2b are hydrogen, R'3 is methyl and R' is pyridyl or cyanophenyl or iiiii) X' is hydrogen, Y' is oxygen, R' is O-chlorine and R'2a and R_{2a} are hydrogen, R'_3 is methyl and R' is phenyl or iiiiii) X'is $-CH_2-CH_2-$, Y' is oxygen, R' is o-chloro, R_{2a} ' and R_{2b} ' are hydrogen, R'3 is methyl and R' is phenyl.